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(54) Title: WATER-DISPERSIBLE OR WATER-SOLUBLE PESTICIDE GRANULES FROM HEAT-ACTIVATED BIN- DERS (57) Abstract <p>Low cost, rapidly water-dispersible or water-soluble granular compositions containing at least 10 % voids and consisting of agglomerates comprised of pesticidal particles bonded together by solid bridges of a water-soluble heat-activated binder, the water-soluble heat activated binder having a melting point range within 40° to 120 °C, a difference of less than 5 °C between the softening point and the onset of solidification, a hydrophile/lipophile balance of about 14 to 19, a dissolution time of not greater than about 50 minutes; and a melt viscosity of at least about 200 cps. Examples of suitable heat-activated binders, which are not intended to be limiting, are ethylene oxide/propylene oxide copolymers and polyethoxyated dinonylphenol.</p>		

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TITLE

5 WATER-DISPERSIBLE OR WATER-SOLUBLE
PESTICIDE GRANULES FROM HEAT-ACTIVATED BINDERS

BACKGROUND OF THE INVENTION

10 In general, water-dispersible or water-soluble granular compositions are prepared by (1) processes involving aqueous (or solvent) spraying and subsequent drying such as pan or fluidized bed granulation, high intensity mixing, granulation, spray drying, or by spraying the active material (or solution thereof) upon a preformed carrier or (2)
15 processes involving compaction such as briquetting, tableting, and extrusion.

Japanese Patent Application No. 52/30577 discloses slow-release agrochemical-urea fertilizers which are formed from liquid or organic solvent
20 solutions of agrochemicals, polyoxyethylene nonionic surfactants, and urea.

U.S. 4,707,287 is directed to the protection of certain enzymes from a peroxyacid bleach granulate and discloses an improved granulate enzyme
25 composition comprising a core of enzyme material and a protective coating comprising an alkaline buffer salt. This patent broadly mentions the term "alkylarylethoxylates" among many others as potential waxy substances used as granulating agents, but there
30 is no teaching of any of the specific heat-activated binders of the instant invention.

SUMMARY OF THE INVENTION

This invention comprises a low cost, rapidly water-dispersible or water-soluble pesticidal
35 granular composition which is comprised of agglomerates consisting essentially of solid

pesticidal particles bound together by solid bridges of a water-soluble heat activated binder (HAB). The granular composition contains at least about 10% voids (preferably 20% or greater) and comprises by weight based on the total weight of the composition:

- (1) 5 to 95% and preferably 20-80% of
pesticidal particles or a mixture of
pesticidal particles having a size in the
range of 1-50 microns or larger if the
pesticide is water-soluble; in
combination with or held together by
solid bridges of
- (2) 5 to 40% and preferably 10-30% of a heat
activated, water-soluble binder (HAB)
having one or more components wherein
said binder meets the following five
criteria:
 - (i) has a melting point range within
40-120°C, and preferably 45 to 100°C;
 - (ii) has a hydrophile/lipophile balance
(HLB) of about 14 to 19, preferably
16-19;
 - (iii) dissolves in mildly-agitated water in
60 min. or less, preferably 50 min. or
less;
 - (iv) has a melt viscosity of at least about
200 centipoise (cps); preferably 1,000
cps or greater, and most preferred
2,000 cps or greater; and
 - (v) has a difference of 5°C, and
preferably 3°C or less between the
softening point and onset of
solidification; and optionally
- (3) one or more additives selected from the
group consisting of

- 5 (i) wicking, physically swelling, or
gas-producing disintegrants;
(ii) anti-caking agents;
(iii) chemical stabilizers;
(iv) co-binders; and
(v) surfactants (wetting agents or
dispersants)

10 such that said composition rapidly forms a high
quality dispersion (or solution) in water, is
attrition resistant (non-dusty), chemically stable,
and non-caking. The agglomerates or granules are
15 150-4000 microns and preferably 250-1500 microns in
size.

DETAILED DESCRIPTION OF THE INVENTION

The most common method of applying agricultural
pesticides involves their dilution in a solvent or
non-solvent liquid in a mix tank followed by spraying
20 of the resulting solution or dispersion. Because of
the increasing costs of non-aqueous solvents and the
toxicity of some of them, formulations involving
water-soluble or water-dispersible granules have
become increasingly popular. In such formulations,
25 the dispersed particles formed on dilution should be
no greater than 50 microns in their largest dimension
to avoid nozzle pluggage or premature settling which
results in uneven application of the pesticide.
Consequently, it is necessary that all of the
30 components of the formulated product rapidly and
completely disperse or dissolve in the dilution water.

Conventional methods for the preparation of
water-soluble or water-dispersible granules involve
(1) solvent spraying such as fluidized bed or pan
35 granulation techniques or the impregnation of an
active pesticide agent on preformed carrier granules
or (2) compaction such as tabletting or extrusion.

Granules prepared by bed or pan granulation are generally sprayable upon dilution with water, while
5 the impregnated or compacted compositions are usually applied dry and mechanically, for example, using spreaders. The solvent spraying processes can yield granules which are rapidly water-dispersible but are expensive, due to the drying step and extensive,
10 space-consuming equipment required. The granules produced from the compaction processes are generally slow in water dispersion. Furthermore, both of these processes often require specialized technology for their operation.

15 Often it is desirable to use mixtures of two or more pesticides of different functions, e.g., a mixture of a herbicide and an insecticide, to provide broad spectrum control over a variety of weeds and/or undesirable organisms. However, some of the
20 individual components are physically or chemically incompatible as mixtures, especially in long-term storage. For example, carbamate insecticides are generally unstable in the presence of alkaline components and sulfonylurea herbicides are known to
25 be unstable in the presence of acidic materials. The chemical incompatibility can be due to an impurity present in the complementary pesticide and not the bioactive component itself. For these reasons it would be desirable to have a sprayable, formulated
30 product consisting of particles or granules wherein potentially incompatible, active components are physically separated.

The present invention comprises low cost, rapidly water-dispersible or water-soluble granular
35 compositions consisting of agglomerates comprised of pesticidal particles bonded together by solid bridges of the heat-activated binder (HAB). The granules

have 10% voids or more, and have a preferred size of 150 to 4,000 microns. The preferred size of
5 pesticidal particles is 1 to 50 microns, especially for pesticides with low water-solubility, to promote water dispersion, avoid premature settling, and avoid nozzle/screen pluggage during tank-mixing or application in the field. Water-soluble pesticidal
10 particles may be larger.

The granular compositions of this invention display a break-up time of three minutes or less in water, good aqueous dispersion properties with a long tube sedimentation value of 0.02 mL or less,
15 attrition of 33% or less, and are preferably non-caking after 100 hours at 45°C under a pressure of 3.5 Kg/cm².

The granules can be comprised of mixtures of pesticidal particles which are ordinarily chemically
20 incompatible (e.g., in a conventional granule made by water spraying, such as fluid bed or pan granulation) because (1) the pesticidal particles can be physically separated from each other via HAB bridges; and (2) water is not required during
25 granulation/drying.

Advantages of the present granules include potential incorporation of incompatible pesticides in the same granule and low cost. The process used to prepare these granules is simple and does not require
30 specialized technology. It utilizes readily available, compact equipment. The process does not require extensive dust collection systems nor a space-consuming and expensive drying operation.

The compositions of this invention can be
35 prepared by several processes (either in a batch or continuous mode) including the processes wherein (1) the pesticidal particles, the HAB particles and

optional particulate additives are tumbled/mixed and heat is applied externally until the granules have grown to the desired size, following which the heat is shut off and the granules are allowed to cool while still tumbling or sitting in a separate container; or where (2) the pesticidal particles, HAB, and optional particulate additives are intensely sheared/mixed such that frictional heat melts the HAB thereby effecting granulation following which the aggregates are then cooled; or where (3) the pesticidal particles and optional particulate additives are tumbled/mixed and are sprayed with the heat-activated binder which has been pre-heated and is in a molten state following which the resulting agglomerates are cooled.

Processes (1) and (3), involving gentle tumbling/mixing, can be carried out, e.g. in a heated fluidized bed, a heated blender (e.g., paddle or ribbon type blenders, vee-blenders, zig-zag blenders, Lodige® blenders, Nauta® mixers) or a heated pan or drum granulator. Process (3) may not require additional heat other than that needed to melt the HAB for spraying. Subsequent cooling of the resulting agglomerates is done either in or outside of the processing vessel. Process (2) involving high intensity mixing/shearing can be carried out e.g., in Schugl® or turbulator-type vessels. In Process (1) a preferred method of preparing the initial mixture of particulates before granulation is to mill the pesticidal active plus additives and then mix (e.g., via tumbling) with HAB particulates (e.g., of a size 500-1,000 microns). Separation of pesticides can be enhanced and incompatibility then reduced (especially when one active is present in minor proportions) by forming granules from a particulate premix of the

major active component, HAB, and additives, followed
by introduction of the minor active component (and
5 optionally additional HAB), while the granules are
not so as to imbed the second active particulates in
a HAB layer on the surface of the first granules.

The term "pesticide" is intended to refer to
biologically active compositions containing chemicals
10 which are effective in killing pests or preventing or
controlling their growth. These chemicals are
commonly known as herbicides, fungicides,
insecticides, nematocides, acaricides, miticides,
virucides, algicides, bactericides, plant growth
15 regulants and their agriculturally suitable salts.
Preferred are those pesticides that have melting
points above 80°C; more preferred are pesticides that
melt above 100°C. The preferred size of the
pesticidal particles used in this invention is 1 to
20 50 microns. Examples of suitable pesticides are
listed below in Table 1.

25

30

35

TABLE 1

HERBICIDES

5	Cmpd.			
	<u>No.</u>	<u>Common Name</u>	<u>M.P. (°C)</u>	<u>Chemical Name</u>
10	1	acifluorfen	142-160	5-[2-chloro-4-(trifluoromethyl)phenoxy]-2-nitrobenzoic acid
15	2	asulam	142-144	methyl [(4-aminophenyl)sulfonyl]carbamate
	3	atrazine	175-177	6-chloro-N-ethyl-N'-(1-methylethyl)-1,3,5-triazine-2,4-diamine
20	4	bensulfuron methyl	185-188	2-[[[(4,6-dimethoxy-2-pyrimidinyl)amino]carbonyl]amino]sulfonyl]methyl]benzoic acid, methyl ester
25	5	bentazon	137-139	3-(1-methylethyl)-(1H)-2,1,3-benzothiadiazin-4(3H)-one, 2,2-dioxide
30	6	bromacil	158-159	5-bromo-6-methyl-3-(1-methylpropyl)-2,4(1H,3H)pyrimidinedione
35	7	bromoxynil	194-195	3,5-dibromo-4-hydroxybenzonitrile

	Cmpd.		m.p. (°C)	Chemical Name
	No.	Common Name		
5	8	chlaramben	200-201	3-amino-2,5-dichlorobenzoic acid
10	9	chlrimuron ethyl	>100	2-[[[(4-chloro-6-methoxy-2-pyrimidinyl)amino]carbonyl]-amino]sulfonyl]benzoic acid, ethyl ester
15	10	chloroxuron	151-152	N'-[4-(4-chlorophenoxy)-phenyl]N,N-dimethylurea
20	11	chlorsulfuron	174-178	2-chloro-N-[[[(4-methoxy-6-methyl-1,3,5-triazin-2-yl)-amino]carbonyl]benzene-sulfonamide
	12	chlortoluron	147-148	N'-(3-chloro-4-methylphenyl)-N,N-dimethylurea
25	13	clomazone	oil	2-[(2-chlorophenyl)methyl]-4,4-dimethyl-3-isoxazolidinone
30	14	cyanazine	166-167	2-[[[4-chloro-6-(ethylamino)-1,3,5-triazin-2-yl]amino]-2-methylpropanenitrile
35	15	dazomet	104-105	tetrahydro-3,5-dimethyl-2H-1,3,5-thiadiazine-2-thione

10

	Cmpd.		M.P. (°C)	Chemical Name
	No.	Common Name		
5	16	desmediphan	120	ethyl [3-[[[(phenylamino)- carbonyl]oxy]phenyl]- carbamate
10	17	dicamba	114-116	3,6-dichloro-2-methoxybenzoic acid
	18	dichlobenil	139-145	2,6-dichlorobenzonitrile
15	19	dichlorprop	117-118	(±)-2-(2,4-dichlorophenoxy)- propanoic acid
	20	diphenamid	134-135	N,N-dimethyl-α-phenylbenzene- acetamide
20	21	dipropetryn	104-106	6-(ethylthio)-N,N'-bis(1- methylethyl)-1,3,5- triazine-2,4-diamine
25	22	diuron	158-159	N'-(3,4-dichlorophenyl)-N,N- dimethylurea
	23	thiameturon	>100	3-[[[(4-methoxy-6-methyl- 1,3,5-triazin-2-yl)amino]- carbonyl]amino]sulfonyl]- 2-thiophenecarboxylic acid, methyl ester
30				
35				

	Cmpd.		m.p. (°C)	Chemical Name
	No.	Common Name		
5	24	----	>100	2-[[[N-(4-methoxy-6-methyl-1,3,5-triazine-2-yl)-N-methylamino]carbonyl]-amino]sulfonyl]benzoic acid, methyl ester
10	25	fenac	156	2,3,6-trichlorobenzeneacetic acid
15	26	fenuron	133-134	N,N-dimethyl-N'-phenylurea
	27	fluometuron	163-164	N,N-dimethyl-N'-[3-(trifluoromethyl)phenyl]urea
20	28	fluridone	151-154	1-methyl-3-phenyl-5-[3-(trifluoromethyl)phenyl]-4(1H)-pyridinone
25	29	fomesafen	220-221	5-[2-chloro-4-(trifluoromethyl)phenoxy]-N-(methylsulfonyl)-2-nitrobenzamide
	30	glyphosate	200	N-(phosphonomethyl)glycine
30	31	hexazinone	115-117	3-cyclohexyl-6-(dimethylamino)-1-methyl-1,3,5-triazine-2,4(1H,3H)-dione
35				

	Compd.		m.p. (°C)	Chemical Name
	No.	Common Name		
5	32	imazamethabenz	>100	6-(4-isopropyl-4-methyl-5-oxo-2-imidazolin-2-yl)-m-toluic acid, methyl ester and 6-(4-isopropyl-4-methyl-5-oxo-2-imidazolin-2-yl)-p-toluic acid, methyl ester
10				
	33	imazaquin	219-222	2-[4,5-dihydro-4-methyl-4-(1-methylethyl)-5-oxo-1H-imidazol-2-yl]-3-quinoline-carboxylic acid
15				
	34	imazethapyr	172-175	(±)-2-[4,5-dihydro-4-methyl-4-(1-methylethyl)-5-oxo-1H-imidazol-2-yl]-5-ethyl-3-pyridinecarboxylic acid
20				
	35	ioxynil	209	4-hydroxy-3,5-diiodobenzo-nitrile
25				
	36	isoproturon	155-156	N-(4-isopropylphenyl)-N',N'-dimethylurea
	37	isouron	119-120	N'-[5-(1,1-dimethylethyl)-3-isoxazolyl]-N,N-dimethylurea
30				
	38	isoxaben	176-179	N-[3-(1-ethyl-1-methylpropyl)-5-isoxazolyl]-2,6-dimethoxybenzamide
35				

13

	Cmpd.		m.p. (°C)	Chemical Name
	No.	Common Name		
5	39	karbutilate	176-178	3-[[[(dimethylamino)carbonyl]-amino]phenyl-(1,1-dimethylethyl)carbamate
10	40	lenacil	316-317	3-cyclohexyl-6,7-dihydro-1H-cyclopentapyrimidine-2,4-(3H,5H)dione
15	41	MCPA	100-115	(4-chloro-2-methylphenoxy)-acetic acid
	42	MCPB	100	4-(4-chloro-2-methylphenoxy)-butanoic acid
20	43	mefluidide	183-185	N-[2,4-dimethyl-5-[[[(trifluoromethyl)sulfonyl]-amino]phenyl]acetamide
25	44	methabenzthiazuron	119-120	1,3-dimethyl-3-(2-benzothiazolyl)urea
	45	methazole	123-124	2-(3,4-dichlorophenyl)-4-methyl-1,2,4-oxadiazolidine-3,5-dione
30	46	metribuzin	125-126	4-amino-6-(1,1-dimethylethyl)-3-(methylthio)-1,2,4-triazin-5(4H)-one
35				

Cmpd.	No.	Common Name	m.p. (°C)	Chemical Name
5	47	metsulfuron methyl	163-166	2-[[[[(4-methoxy-6-methyl- 1,3,5-triazin-2-yl)amino]- carbonyl]amino]sulfonyl]- benzoic acid, methyl ester
10	48	monuron	174-175	N'-(4-chlorophenyl)-N,N- dimethylurea
15	49	naptalam	185	2-[(1-naphthalenylamino)- carbonyl]benzoic acid
	50	neburon	102-103	1-butyl-3-(3,4-dichloro- phenyl)-1-methylurea
20	51	nitralin	151-152	4-(methylsulfonyl)-2,6- dinitro-N,N-dipropyl- aniline
25	52	norflurazon	174-180	4-chloro-5-(methylamino)-2- [3-(trifluoromethyl)phenyl]- 3(2H)-pyridazinone
30	53	oryzalin	141-142	4-(dipropylamino)-3,5-dinitro- benzenesulfonamide
	54	perfluidone	142-144	1,1,1-trifluoro-N-[2-methyl- 4-(phenylsulfonyl)phenyl]- methanesulfonamide
35	55	phenmedipham	143-144	3-[(methoxycarbonyl)amino]- phenyl (3-methylphenyl)- carbamate

15

Cmpd.			
No.	Common Name	m.p. (°C)	Chemical Name
5			
56	picloram	>215 (DEC)	4-amino-3,5,6-trichloro-2-pyridinecarboxylic acid
10			
57	prometryn	118-120	N,N'-bis(1-methylethyl)-6-(methylthio)-1,3,5-triazine-2,4-diamine
58	pronamide	155-156	3,5-dichloro-N-(1,1-dimethyl-2-propynyl)benzamide
15			
59	propazine	212-214	6-chloro-N,N'-bis(1-methylethyl)-1,3,5-triazine-2,4-diamine
20			
60	pyrazon	205-206	5-amino-4-chloro-2-phenyl-3(2H)pyridazinone
61	siduron	133-138	N-(2-methylcyclohexyl)-N'-phenylurea
25			
62	simazine	225-227	6-chloro-N,N'-diethyl-1,3,5-triazine-2,4-diamine
63	sulfometuron methyl	182-189	2-[[[(4,6-dimethyl-2-pyrimidinyl)amino]carbonyl]-amino]sulfonyl]benzoic acid, methyl ester
30			
64	tebuthiuron	161-164	N-[5-(1,1-dimethylethyl)-1,3,4-thiadiazol-2-yl]-N,N'-dimethylurea
35			

16

Cmpd.				
No.	Common Name	M.P. (°C)	Chemical Name	
5	65	terbacil	175-177	5-chloro-3-(1,1-dimethyl-ethyl)-6-methyl-2,4(1H,3H)-pyrimidinedione
10	66	terbutyl- azine	177-179	2-(tert-butylamino)-4-chloro-6-(ethyl-amino)-s-triazine
	67	terbutryn	104-105	N-(1,1-dimethylethyl)-N'-ethyl-6-(methylthio)-1,3,5-triazine-2,4-diamine
15	68	triclopyr	148-150	[(3,5,6-trichloro-2-pyridinyl)oxy]acetic acid
20	69	2,4-D	140	(2,4-dichlorophenoxy)acetic acid
	70	2,4-DB	119-120	4-(2,4-dichlorophenoxy)-butanoic acid
25	71	triasulfuron	>100	(3-(6-methoxy-4-methyl-1,3,5-triazin-2-yl)-1-[2-(2-chloroethoxy)phenylsulfonyl]urea
30	72	primisulfuron	>100	[2-/3-(4,6-bis(difluoromethoxy)pyrimidin-2-yl-ureidosulfonyl)benzoic acid methylester]
35				

17

Cmpd.				
No.	Common Name	m.p.(°C)	Chemical Name	
5	73	----	>100	[2-/3-(4,6-bis(difluoro-methoxy)-pyrimidin-2-yl)-ureidosulfonyl]-benzoic acid methylester]
10	74	NC-311	170-172	[5-pyrazolesulfonamide, N-[(4-methoxy-6-methyl-pyrimidine-2-yl)-amino-carbonyl]-4-methoxy-carbonyl-1-methyl-]
15	75	----	160-162	N-[[[(4,6-dimethoxy-2-pyrimidinyl)amino]carbonyl]-3-(ethylsulfonyl)-2-pyridinesulfonamide
20	76	----	152-159	2-[[[(4,6-dimethoxy-2-pyrimidinyl)amino]carbonyl]-amino]sulfonyl]-N,N-dimethyl-3-pyridine-carboxamide
25	77	----	204-206	Methyl 2-[[[(4-ethoxy-6-(methylamino)-1,3,5-triazin-2-yl)amino]carbonyl]amino]-sulfonyl]benzoate
30				
<u>FUNGICIDES</u>				
35	78	carbendazim	302-307	methyl 2-benzimidazole-carbamate

18				
	<u>Cmpd.</u>			
	<u>No.</u>	<u>Common Name</u>	<u>M.P. (°C)</u>	<u>Chemical Name</u>
5	79	thiuram	146	tetramethylthiuram disulfide
	80	dodine	136	n-dodecylguanidine acetate
10	81	chloroneb	133-135	1,4-dichloro-2,5-dimethoxy- benzene
	82	cymoxanil	160-161	2-cyano-N-ethylcarbamoyl- 2-methoxyiminoacetamide
15	83	captan	178	N-trichloromethylthiotetra- hydrophthalamide
	84	folpet	177	N-trichloromethylthio- phthalimide
20	85	thiophanate- methyl	195	dimethyl 4,4'-(o-phenylene)- bis(3-thiocallophanate)
25	86	thiabendazole	304-305	2-(thiazol-4-yl)benzimidaz- ole
	87	chlorothalonil	240-241	tetrachloroisophthalo- nitrile
30	88	dichloran	195	2,6-dichloro-4-nitroaniline
	89	captafol	160-161	cis-N-[1,1,2,2-tetrachloro- ethylthio)cyclohex-4- ene-1,2- dicarbioximide
35				

	Cmpd.		m.p. (°C)	Chemical Name
	No.	Common Name		
5	90	iprodione	133-136	3-(3,5-dichlorophenyl)-N-(1-methylethyl)-2,4-dioxo-1-imidazolidine carboxamide
10	91	vinclosolin	108	3-(3,5-dichlorophenyl)-5-ethenyl-5-methyl-2,4-oxazolidinedione
15	92	kasugamycin	202-204 (DEC)	kasugamycin
20	93	triadimenol	121-127	beta-(4-chlorophenoxy)-alpha-(1,1-dimethylethyl)-1H-1,2,4-triazol-1-ethanol
25	94	flutriafol	130	+alpha-(2-fluorophenyl)-alpha-(4-fluorophenyl)-1H-1,2,4-triazole-1-ethanol
25	95	flusilazol	52-53 HCl 201-203	1-[[bis(4-fluorophenyl)-methylsilyl)methyl]-1H-1,2,4-triazole
30	96	hexaconazole	111	(+/-)-alpha-butyl-alpha-(2,4-dichlorophenyl)-1H-1,2,4-triazole-1-ethanol
35	97	fenarimol	117-119	alpha-(2-chlorophenyl)-alpha(4-chlorophenyl)-5-pyridinemethanol

Cmpd.

	No.	Common Name	m.p. (°C)	Chemical Name
5				
		BACTERICIDES		
10	98	oxytetracycline dihydrate	181-182 (DEC)	oxytetracycline dihydrate
		ACARICIDES		
15	99	hexathiasox	108-109	trans-5-(4-chlorophenyl)-N-cyclohexyl-4-methyl-2-oxo-3-thiazolidinecarboxamide
20	100	oxythioquinox	169-170	6-methyl-1,3-dithiolo-[2,3-B]quinonolin-2-one
	101	dienochlor	122-123	bis(pentachloro-2,4-cyclopentadien-1-yl)
25	102	cyhexatin	245	tricyclohexyltin hydroxide
		INSECTICIDES		
30	103	carbofuran	150-152	methylcarbamic acid, ester with 2,3-dihydro-2,2-dimethyl-7-benzofuranol
35	104	carbaryl	142	methylcarbamic acid, ester with 8-naphthol

21

Cmpd.	No.	Common Name	m.p. (°C)	Chemical Name
5	105	thiodicarb	173-174	dimethyl N,N'-[thiobis-(N-methylimino)carbonyl-oxy]]-bis[ethanimido-thioate]
10	106	deltamethrin	98-101	α -cyano-3-phenoxybenzyl-cis-3-(2,2-dibromovinyl)-2,2-dimethylcyclopropane carboxylate
15				
20				
25				
30				
35				

The term "heat-activated binder" refers to any surface active material comprised of one or more components which dissolve rapidly in water, have some viscosity near the melting point for tackiness, and are thus capable of acting as a binder when heat is applied. At some elevated temperature, the binder softens and melts, thereby becoming sticky enough to bind the pesticidal particles into granules. A more preferred amount of binder used in this invention is 10-30% by weight based on the total weight of the composition. A more preferred melting point range for the binders of this invention is 45°C to 100°C. Examples of suitable heat-activated binders, which are not intended to be limiting, are ethylene oxide/propylene oxide copolymers and polyethoxylated dinonylphenol.

The HAB can be a single component or multi-components which are mixed in the solid state, co-melted or co-dissolved. Preferred single component HAB's are ethylene oxide/propylene oxide copolymers and polyethoxylated dinonylphenol. Specifically preferred single components are block copolymers of ethylene oxide/propylene oxide, where 80% is ethylene oxide and 20% is propylene oxide, and polyethoxylated dinonylphenol with 150 ethylene oxide units. The preferred copolymer has an HLB of 16 and a melting point of about 45 to 61°C. The preferred dinonylphenol derivative has an HLB of about 19 and a melting point of about 48 to 63°C.

The HAB must meet the following five criteria:

- (1) have a melting point range within 40 to 120°C;
- (2) be water-soluble with a hydrophile/lipophile balance (HLB) of about 14 to 19;

(3) dissolve in mildly-agitated water in 50 minutes or less;

5 (4) have a melt viscosity of at least 200 cps; and

(5) have a difference of 5°C or less between the softening point and onset of solidification.

10 The use of a HAB having a very low melting point can lead to caking of the granules, while use of a HAB having a very high melting point can require a temperature sufficiently high so that decomposition of the pesticide or other components can occur during granulation.

15 Surface activity as measured by the critical HLB range is necessary to provide good bonding of the HAB to pesticidal particles and rapid wetting at the onset of bridge dissolution when the granules are placed in water. Materials which have too low an HLB
20 are not completely water-soluble.

The dissolution rate in water is very important, since factors other than HLB affect dissolution, e.g., viscosity of the hydrated HAB and its tendency to form a gel-like layer when in contact
25 with mildly or non-agitated water.

The use of a HAB having the specified melt viscosity and minimum difference between softening and solidification temperatures is necessary so that it will be tacky enough to effect agglomeration of
30 pesticidal particles near the melting point of the HAB.

Additives, many of which are commonly used in conventional granules, may optionally be used in HAB granules. Examples include:

35 (1) disintegrants which wick in water, physically expand, or produce gas to aid break-up of

the granule. Non-limiting examples of suitable disintegrants include cross-linked polyvinyl pyrrolidone, microcrystalline cellulose, cross-linked sodium carboxymethyl cellulose, salts of polyacrylates of methacrylates, and the combination of sodium or potassium bicarbonates or carbonates with acids such as citric or fumaric acid, used alone or in combination, at levels of up to 30% by weight based on the total weight of the composition;

(2) anticaking agents to prevent clumping of granules when stored under hot warehouse conditions. Non-limiting examples of suitable anticaking agents include sodium or ammonium phosphates, sodium carbonate or bicarbonate, sodium acetate, sodium metasilicate, magnesium or zinc sulfates, magnesium hydroxide (all optionally as hydrates), and sodium alkylsulfosuccinates;

(3) chemical stabilizers to prevent decomposition of the active(s) during storage. Non-limiting examples of suitable chemical stabilizers include alkaline earth or transition metal sulfates such as magnesium, zinc, aluminum, and iron (optionally as hydrates) used at levels of 1-9% by weight based on the total weight of the composition;

(4) co-binders to achieve optimized properties such as increased granulation efficiency or improved anticaking. Up to 50% co-binders such as polyethylene glycols, polyethylene oxide, polyethoxylated fatty acids or alcohols, hydrated inorganics such as sodium silicate, sorbitol, or urea may be used; and

(5) surfactants to improve the speed and quality of wetting and dispersion of the granule upon

mixing with water. Often dispersing agents are most useful, since the HAB itself has wetting characteristics.

Examples of preferred dispersants include sodium or ammonium salts of sulfonated naphthalene (or methyl naphthalene)-formaldehyde condensates, sodium, calcium, or ammonium salts of ligninsulfonates (optionally polyethoxylated); dialkyl; diolalkynes; sodium taurates; and sodium or ammonium salts of maleic anhydride copolymers.

HAB candidates may be identified by the following tests:

(1) the melting point is measured by DSC (Differential Scanning Calorimetry) at a 5°C/minute heating rate. The onset of the melt should be no lower than 40°C;

(2) The hydrophile/lipophile balance with a total possible range of 1 to 20 is determined by the method outlined in McCutcheon's "Detergents and Emulsifiers", 1971 annual, page 223;

(3) The rate of dissolution in water is determined by the following procedure:

- (a) a sample of the test material (0.15 g) is placed in the bottom of a glass graduated cylinder with an inside diameter of 2.8 cm,
- (b) the cylinder is placed on a steam bath (alternately heated externally with a stream of hot air near the bottom) until the sample is fully melted,
- (c) the cylinder is placed on a level surface and the sample allowed to

solidify upon cooling to 25°C,
yielding an even layer in the
bottom,

(d) water is added (100 mL at 25°C) to
the cylinder and is stirred at 110
rpm with a rectangular metal or
plastic paddle having a thickness
of 1.5 mm, a width of 18 mm, and a
height of 16.5 mm so that the
bottom of the paddle is 48 cm above
the surface of the solidified
sample, and

(e) the time for complete dissolution
of the sample is noted;

(4) The viscosity at the softening point is
calculated using an Arrhenius plot (\ln viscosity vs
 $1/T$).

The plot is derived from experimental viscosity
measurements vs temperature using a rotational
viscometer operated at a shear rate of 1.16 sec^{-1} .
Viscosity measurements are taken over at least a 30°C
temperature range whose minimum temperature is within
1°C of the softening point as measured by DSC.

Another requirement of the melt behavior of HAB
candidates is that the onset of softening in the
heating curve differs by 5°C or less from the onset
of solidification in the consecutive cooling curve.

This parameter is measured using a differential
scanning calorimeter (e.g., Du Pont Instruments 1090
Thermal Analyser with model 910 DSC module). One to
three milligrams of sample is typically used in a
hermetically-sealed coated aluminum pan. The heating
curve endotherm is observed at 5°C/minute, while the
cooling curve exotherm is observed at 1°C/minute.

Typically a sample is heated from 25°C to 100°C to 120°C and then allowed to cool back to 25°C. It should be noted that a given HAB can exhibit a broad melting behavior (usually 12 to 16°C from softening to full melt).

Granules of this invention have at least 10% voids and preferably at least 20%. The upper limit of voids is set by the fragility (high attrition) of the granule. Determination of voidage is accomplished by pycnometer measurements of the starting premix powder and the final HAB granules, using a paraffin oil. Alternately, helium porosimetry may be used. Voids are important to speed penetration of water into the granule and thus aid break-up in the mix tank.

The granules also exhibit break-up times in water of less than 3 minutes and preferably less than 2. Break-up time is measured by adding a sample of the granules (0.5 g, 250 to 1410 microns) to a 100 mL graduated cylinder [internal height after stoppering is 22.5 cm; I.D. is 28 mm] containing 90 mL of distilled water at 25°C, following which the cylinder is clamped in the center, stoppered, and rotated about the center at 8 rpm until the sample is completely broken up in the water.

Formation of a high quality aqueous dispersion is also a desirable property and is determined by the long tube sedimentation test in U.S. 3,920,442 (Col. 9, lines 1 to 39). Acceptable values correspond to 0.02 mL, preferably 0.01 mL of solids after 5 minutes of settling.

The granules should exhibit low attrition characteristics which can be determined by the attrition test in U.S. 3,920,442 (Col. 8, lines 5 to 48). The test is modified to use test samples of the

commercial granule size (e.g., 250 to 1410 microns). Attrition values of less than 40% and preferably less than 30% are acceptable.

The granules should also resist caking. This property is determined by taping a stainless steel disc (0.9 mm thick x 51 mm diameter) flush with the bottom of a glass cylinder (46.5 mm i.d. x 75 mm length x 51 mm thickness) following which the granular sample (20 g) is delivered to the cylinder assembly and leveled, and a second stainless steel disc (0.9 mm thick x 44.5 mm diameter) is placed on the top of the granules.

A 400 g weight (45 mm diameter or less) is then placed on top of the inner disc, and the entire assembly is placed in an oven for 100 hours at 45°C (preferably 55°C) following which the assembly is removed from the oven, the weight removed, and the sample allowed to cool to room temperature. The bottom disc is then detached and if the sample flows out of the cylinder, the resistance to caking is excellent, and if the sample remains in the cylinder, the cake is removed, placed onto a flat surface and a penetrometer is used with a single-edged razor to measure the minimum force necessary to cleave the cake.

Cakes requiring a force of less than 100 g, and preferably less than 5 g are acceptable.

The following examples are presented to illustrate, but not to restrict, this invention.

Definitions of Ingredients Used in Examples

	<u>Name</u>	<u>Identity</u>
5	Macol® DNP 150 (Mazer Chemicals)	polyethylated dinonylphenol (150 ethylene oxide units)
10		<u>Melt behavior:</u> <ul style="list-style-type: none">• melting point - softening point 48°C; finish 63°C• difference between softening point and onset of solidification = 2°C
15		<ul style="list-style-type: none">• melt viscosity - about 1,900 cps at softening point
20		<u>Dissolution Rate:</u> 19 minutes
		<u>HLB:</u> 19
25	Pluronic® F108 (BASF)	ethylene oxide/propylene oxide block copolymer with 80% ethylene oxide and 20% propylene oxide units
30		<u>Melt behavior:</u> <ul style="list-style-type: none">• melting point - softening point 45°C; finish 61°C• difference between softening point and onset of solidification = 0°C
35		

30

	<u>Name</u>	<u>Identity</u>
5		<ul style="list-style-type: none">• melt viscosity: at softening point about 26,500 centipoises (cps)
10		<u>Dissolution Rate:</u> 50 minutes <u>HLB:</u> 16
15	Hodag® E100 (Hodag Chemical Corp.)	a 100 mole ethoxylate of nonylphenol <u>Melt behavior:</u> <ul style="list-style-type: none">• melting point - softening point 40°C; finish point 64°C• difference between softening point and onset of solidification = 0°C• melt viscosity at softening point = 1,100 cps
20		
25		<u>Dissolution Rate:</u> 20 minutes
30		<u>HLB:</u> 19

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	<u>Name</u>	<u>Identity</u>
5	Iconol® OP-40 (BASF)	a 40 mole ethoxylate of octylphenol
		<u>Melt behavior:</u>
10		• melting point - softening point = 40°C; finish = 55°C
		• difference between softening point and onset of solidification = 3°C
15		• viscosity at softening point about 700 cps
		<u>Dissolution Rate:</u> 18 minutes
20		<u>HLB:</u> 18
	Polyplasdone® XL-10 (GAF)	Cross-linked polyvinyl pyrrolidone
25	Avicel® PH-105 (FMC)	microcrystalline cellulose
	AC-DI-SOL® (FMC)	Cross-linked sodium carboxymethyl cellulose
30	Morwet® D425 (Desoto)	sodium naphthalene sulfonate formaldehyde condensate
35	Morwet® EFW (Desoto)	sodium alkyl naphthalene sulfonate

	<u>Name</u>	<u>Identity</u>
5	Lignosol® TSF (Reed)	ammonium lignosulfonate
	Monawet® MB100 (Mona)	sodium dibutylsulfosuccinate
10	Aerosol® A196 (Amer. Cy.)	sodium dicyclohexylsulfo- succinate (+ 15% sodium benzoate)
15	Explotab® (Edward Mendell Co.)	sodium starch glycolate
	Triton® AG-120 (R & H)	polyethoxylated nonyl phenol absorbed on silica
20	Triton® X-120 (R & H)	polyethoxylaed nonyl phenol absorbed in MgCO ₃

EXAMPLE 1

- 25 All ingredients below (with the exception of
the Macol®) were mixed and then milled on a high
intensity rotary shearing mill. The resultant
mixture was then mixed with Macol® DNP 150 (<840
microns) to form a premix for granulation. A 150 g
30 portion of the premix was placed in a fluidized bed
and heated air was applied to the granules. When the
temperature of the granules reached 70°C (about 12
minutes) the heat was shut off and the granules
allowed to cool while still fluidized by unheated
35 air. A 70% yield of a 250 to 1410 micron spherical
granules was realized. The premix formulation and
resulting properties of the granules are given below.

Premix Formulation

5		<u>Weight Percent</u>
	Chlorsulfuron	75.0
	Macol® DNP150	10.0
	Anhydrous MgSO ₄	6.0
	Polyplasdane® XL-10 +	9.0
10	Impurities	

Properties of Granules (250 to 1410 microns)

	% Attrition	21.0%
	25°C Break-up Time	
15	in water	75.0 sec
	OC Break-up Time	
	in water	102.0 sec
	25°C Break-up Time -	
	in 28-0-0 Liquid	
20	Fertilizer	214.0 sec
	55°C Caking	0.0 g
	Long Tube	
	Sedimentation	Trace
	Long Tube	
25	Sedimentation	
	(1 wk/55°C)	.003
	Assay (% chlor-	
	sulfuron) Control	72.9%
	1 wk/55°C	70.0%
30		

EXAMPLE 2

Example 1 was repeated except that the premix without binder was hammer-milled.

	<u>Premix Formulation</u>	<u>Example 2</u>
5	Chlorsulfuron Tech (%)	77
	Macol® DNP150 (%)	10
	MgSO ₄ ·7H ₂ O (%)	3
	ZnSO ₄ ·7H ₂ O (%)	3
	Polyplasdone® XL-10 (%)	7
10	Avicel® PH-105 (%)	-
	AC-DI-SOL® (%)	-
	Premix Charge (GM)	1934
	Conversion (%)	73
	Attrition (%)	29
15	Long tube sedimentation	0.002
	Long tube sedimentation (1 wk/55°C)	0.002
	25°C H ₂ O Break-up (sec)	83
	0°C H ₂ O Break-up (sec)	107
20	25°C 28-0- Break-up (sec)	250
	55°C Caking (GM Force)	0

EXAMPLE 3

Approximately 73.84 g of 2,4-D, Na salt and
 25 1.16 g of 2-[[N-(4-methoxy-6-methyl-1,3,5-triazin-
 2-yl)-N-methylamino]carbonylamino]sulfonyl]benzoic
 acid, methyl ester, Na salt were milled together as
 in Example 1. This material was then blended with 25
 g of Macol® DNP-150 (<840 microns). The mixture was
 30 then added to a laboratory double cone blender and
 heated with a heat gun to 77°C, whereupon granulation
 was observed. The heat was removed and the granules
 allowed to cool to 50°C then removed from the
 blender. Approximately 97.3 g were recovered with
 35 88.3% being between 250 and 1410 microns in size.
 The physical properties of the granules were: long
 tube sedimentation (5 minute reading) 0 ml,

attrition-33.7%, break-up time in 25°C water-2.17 minutes and bulk density-0.50 g/ml.

- 5 The granules exhibited good chemical stability upon aging, also with no loss of the above physical properties.

EXAMPLE 4

- 10 Approximately 72.86 g 2,4-D, Na salt, 1.14 g of 2-[[N-(4-methoxy-6-methyl-1,3,5-triazin-2-yl)-N-methylamino]carbonyl]amino]sulfonyl]benzoic acid, methyl ester, Na salt and 1 g NaHCO₃ were milled together as in Example 1. This mixture was then blended with 25 g Macol® DNP-150 (<840 microns). The
15 procedure of Example 1 was then followed to produce granules. Approximately 93.4 g were recovered with 90.9% being between 250 and 1410 microns. The physical properties of the as made granules were: long tube sedimentation (5 minute reading)-trace,
20 bulk density-0.5 g/ml, attrition 37.5%, and break-up time in 25°C water-2.18 minutes. The properties after aging at 45°C for 3 weeks were: long tube sedimentation (5 minute reading)-trace, bulk density-0.5 g/ml, attrition-36.8%, break-up time in
25 25°C water-2.19 minutes. As in Example 3, these granules also maintained good chemical stability on aging.

EXAMPLE 5

- 30 Approximately 1480 g methabenzthiazuron, 9.80 g ammonium salt of chlorsulfuron (technical), 5.16 g Sellogen® HR, 6.88 g Petro® D425, 12.90 g diammonium hydrogen phosphate, 137.26 g kaolin clay, 40 g MgSO₄, and 48 g Polyplasdane® XL-10 were milled in an ACM
35 mill at 90 g/min feed rate, rotor speed of 11,000 rpms, classifier speed of 6,000 rpms and as air flow of 50 cfm. Approximately 1643 g of milled material

was recovered from the mill. Three 600 g batches were granulated in a laboratory 2.2 liter vee blender by combining 522 g of the milled material and 78 g of less than 500 micron Pluronic® F108. This material was blended and heated as in Example 3 to 70°C when granulation was observed. The heat was removed and the granules cooled to 45°C before removing from the blender. Approximately 1787 g of granules were recovered from the blender with 88.7% being between 250 to 1410 microns in size. The physical properties of the granules were: long tube sedimentation (5 minute reading)-0.015 ml, break-up time in 25°C water-1.70 minutes, bulk density-0.5 g/ml, and attrition-11.9%. This material passed both the 45°C and 55°C caking test.

EXAMPLE 6

The following were hammer-milled:
86.9% Na 2,4-D tech (83% assay as acid)
1.3% 2-[[N-(4-methoxy-6-methyl-1,3,5-triazine-2-yl)-N-methylamino]carbonyl]-amino]sulfonyl]benzoic acid, methyl ester,
Na salt (92% assay as the free sulfonylurea)
5.9% Morwet® D42
5.9% Morwet® EFW

The resulting premix was continuously auger fed [62 g/min.] to a 35.6 cm diameter disc agglomerator (56° angle with horizontal) rotating at 30 rpm. Molten Macol® DNP-150 (95°C) was sprayed continuously (23 g/minute) onto the premix in the agglomerator using an external mix, air-atomizing, spray nozzle. The Macol® comprised 25-30% of the total mass of resulting granules. A yield of 61% of 1410 to 1680 micron granules was obtained. The granules had an attrition of 39%, a long tube sedimentation of

0 mL, and a break-up time in 25°C water of 150 seconds.

5

EXAMPLE 7

The premix of Example 1 was metered continuously to a 2 liter capacity stainless steel drum (10 cm high x 12 cm diameter) which was rotated at 34 rpm at a 30° angle with the horizontal. The bed of premix on the drum was maintained at 70-77°C by heating the external wall of the drum with an infrared lamp. Approximately 89% of granules exiting the drum were 250 to 1410 microns in size. These granules had a long tube sedimentation of 0.01 ml, an attrition of 40%, and a break-up time of 64 seconds in 25°C water.

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EXAMPLE 8

A premix of 20 g of sodium 2,4D (84% assay), 0.5 g 2-[[N-(4-methoxy-6-methyl-1,3,5-triazin-2-yl)-N-methylamino]carbonyl]amino]sulfonyl]benzoic acid, methyl ester, Na salt (91% assay), and 3.6 g of Maco10 DNP-150 were milled as in Example 1 for two minutes. The dustless contents were then screened, revealing that 61% of the granules produced were in the 149 to 840 micron range and 89% in the 74 to 840 micron range. The long tube sedimentation of the granules in the latter size range was 0 ml, the break-up time in 25°C water was 90 seconds, and the attrition was 40%. There was 0% decomposition of either active after aging 1 week and 3% decomposition of the sulfonylurea after 2 weeks at 55°C.

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EXAMPLE 9

Approximately 100 g of premix was made by
5 combining the following ingredients:

	Chlorsulfuron technical	78.5 g
	Macol® DNP-150	12.0 g
	ZnSO ₄ ·7H ₂ O	2.0 g
	MgSO ₄ ·7H ₂ O	2.0 g
10	Ac-Di-Sol®	2.75 g
	Avicel® PH-105	2.75 g

The premix was milled as in Example 1 to a
powder which was then placed in a fluid bed
15 granulator and fluidized with hot air. The bed was
gradually heated to 70°C (9-10 minutes). The
granules formed as the binder softened. The heat was
removed and the granules were allowed to cool while
fluidization was continued. After cooling, the
20 granulated product was sieved. Approximately 76 g of
granules were recovered in the 250 to 1410 micron
size range which exhibited the following properties.

	Break-up Time in 25°C water	81 seconds
25	Caking (1 day/55°C/3.5 Kg/cm ²)	none
	Assay on sample stored 1 week at -6°C	74.4%
	Assay on sample stored 1 week at 55°C	75.8%
30	Long Tube Sedimentation (before and after aging)	0.005 mL
	Attrition	12%

EXAMPLE 10

100 g of a premix was prepared by combining:

5	Chlorsulfuron technical	77.0 g
	Hodag® E-100	10.0 g
	Anhydrous sodium carbonate	2.0 g
	Polyplasdone® XL-10	2.0 g
10	Sodium acetate trihydrate	2.75 g

The premix was milled and granulated as described in Example 9. Approximately 60 g of 250 to 1410 micron granules were recovered. Break-up times in 25°C water averaged 91 seconds. There was no caking after 4 days at 55°C.

EXAMPLE 11

100 g of a premix was prepared by combining:

20	Chlorsulfuron technical	77.0 g
	Iconol® OP-40	10.0 g
	Anhydrous sodium carbonate	6.0 g
	Polyplasdone® XL-10	7.0 g
	Sodium acetate trihydrate	2.75 g

25 The premix was milled and granulated as described in Example 9. Approximately 57 g of 250 to 1410 micron granules were obtained. The break-up time in 25°C water was 69 seconds. There was no caking after 4 days at 55°C. Attrition was 34%.

30

EXAMPLE 12

100 g of a premix was prepared by combining:

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5	Benzoic Acid, 2-[[[4-ethoxy-6-	
	methyldamino-1,3,5-triazin-	
	2-yl)aminocarbonyl]amino-	
	sulfonyl]methyl ester	77.0 g
	Hodag® E-100	10.0 g
10	Anhydrous MgSO ₄	6.0 g
	Polyplasdone® XL-10	7.0 g

The premix was milled and granulated as described in Example 9. Approximately 59 g of 250 to 1410 micron granules were obtained. The break-up time in 25°C water was 90 seconds. The granules did not cake after 4 days at 55°C. Attrition was 28% and long tube sedimentation was 0.005 ml. Chemical stability was excellent.

EXAMPLE 13

The granulation procedure of Example 5 was repeated, using the following ingredients in the premix:

25	methabenzthiazuron	1460 g
	Chlorsulfuron tech.	10 g
	Pluronic® F108	240 g
	MgSO ₄	10 g
	ZnSO ₄ ·7H ₂ O	90 g
30	Morwet® D-425	50 g
	Avicel® PH-105	140 g

The resulting granules (250 to 1410 microns) were produced in 83% yield and had the following properties: long tube sedimentation-0.015 ml, caking-100 g at 45°C, break-up time in 25°C water-90 seconds, and attrition-10%.

CLAIMS

What is claimed is:

5

1. A granular composition which comprises by weight based on the total weight of the composition: 5 to 95% of pesticidal particles in combination with 5 to 40% of a water-soluble heat activated binder having a melting point range within 40° to 120°C, a difference of less than 5°C between the softening point and the onset of solidification, a hydrophile/lipophile balance of about 14 to 19, a dissolution time of not greater than about 50 minutes; and a melt viscosity of at least about 200 cps; and optionally at least one additive selected from the group consisting of

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- (i) wicking, physically swelling, or gas-producing disintegrants;
- (ii) anti-caking agents; and
- (iii) chemical stabilizers; and
- (iv) surfactants (wetting or dispersants) agents and mixtures of the foregoing.

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2. A composition of Claim 1 comprising by total weight 20 to 80% of pesticidal particles, and 10 to 30% of the heat activated binder.

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3. The composition of Claim 1 wherein the difference between the softening point and the onset of solidification is less than 3°C.

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4. The composition of Claim 2 wherein the difference between the softening point and the onset of solidification is less than 3°C.

5. The composition of Claim 1 wherein the melting point range is 45-100°C.

5

6. The composition of Claim 2 wherein the melting point range is 45-100°C.

7. The composition of Claim 3 wherein the melting point range is 45-100°C.

10

8. The composition of Claim 2 wherein the hydrophile/lipophile balance is in the range of 16-19.

15

9. Water-dispersible or water-soluble pesticidal granules which contain at least about 10% voids and comprise agglomerates having a size in the range 150 to 4,000 microns which agglomerates are comprised of pesticidal particles having a size in the range of 1 to 50 microns in diameter bonded together by solid bridges of a water-soluble heat-activated binder as described in Claim 1.

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10. The granules of Claim 9 which contain at least about 20% voids.

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11. The granules of Claim 10 which contain at least about 20% voids and the binder of Claim 2.

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12. The granules of Claim 11 which contain the binder of Claim 8.

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13. The composition of Claim 1 wherein the binder is selected from the class consisting of polyethoxylated dinonylphenol, ethylene oxide/-propylene oxide copolymer and mixtures of the foregoing.


14. The composition of Claim 8 wherein the binder is selected from the class consisting of polyethoxylated dinonylphenol, ethylene oxide/-propylene oxide copolymer and mixtures of the foregoing.

15. The composition of Claim 9 wherein the binder is selected from the class consisting of polyethoxylated dinonylphenol, ethylene oxide/-propylene oxide copolymer and mixtures of the foregoing.

16. The composition of Claim 12 wherein the binder is selected from the class consisting of polyethoxylated dinonylphenol, ethylene oxide/-propylene oxide copolymer and mixtures of the foregoing.

INTERNATIONAL SEARCH REPORT

International Application No PCT/US 91/01105

I. CLASSIFICATION OF SUBJECT MATTER (If several classification symbols apply, indicate all) *		
According to International Patent Classification (IPC) or to both National Classification and IPC		
IPC ⁵ : A 01 N 25/12, 25/14		
II. FIELDS SEARCHED		
Minimum Documentation Searched *		
Classification System	Classification Symbols	
IPC ⁵	A 01 N	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched *		
III. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of Document, ** with indication, where appropriate, of the relevant passages **	Relevant to Claim No. **
X	EP, A, 0206537 (STAUFFER CHEMICAL CO.) 30 December 1986 see page 6, line 16 - page 7, line 24; table 1; page 11, line 32 - page 12, line 5; examples; table III; claims ---	1-8
X	EP, A, 0256608 (STAUFFER CHEMICAL CO.) 24 February 1988 see page 4, lines 20-55; table 1; page 8, lines 20-53; examples; claims ---	1-8
P, X	FR, A, 2645709 (SUMITOMO CHEMICAL CO.) 19 October 1990 see page 2, lines 11-32; claims ---	1
X	Derwent Central Patents Index, Basic Abstracts Journal, Section C, AGDOC, week T37, Derwent Publications Ltd, (London, GB), & JP, B, 52030577 (SUMITOMO CHEMICAL CO. LTD) 9 August 1977 see CO3 no. 59251T/37 cited in the application -----	1
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IV. CERTIFICATION		
Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report	
31st May 1991	25. 07. 91	
International Searching Authority	Signature of Authorized Officer	
EUROPEAN PATENT OFFICE	F.W. HECK 	

**ANNEX TO THE INTERNATIONAL SEARCH REPORT
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US 9101105
SA 45041

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Patent document cited in search report	Publication date	Patent family number(s)	Publication date
EP-A- 0206537	30-12-86	AU-B- 590570	09-11-89
		AU-A- 5801486	04-12-86
		CA-A- 1268639	08-05-90
		JP-A- 61277603	08-12-86
		US-A- 4931080	05-06-90
EP-A- 0256608	24-02-88	JP-A- 63051307	04-03-88
FR-A- 2645709	19-10-90	JP-A- 2279604	15-11-90
		AU-A- 5296690	25-10-90
		GB-A- 2230954	07-11-90
		SE-A- 9001363	19-10-90

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